PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Application No.:	10/635,971	Confirmation No.: 1239
Applicant:	Watanabe *et al.*	
Filed:	August 6, 2003	
Title:	Processes for Preparing 1,3-Dioxolane Nucleosides	
TC/A.AU.:	1644	
Examiner:	Unassigned	
Docket No.:	08841.105052 (PHA 2030 US)	
Customer No.:	20786	

Commissioner for Patents
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Respectfully submitted,

Madeline I. Johnston, Esq.
Reg. No. 36,174

Date: February 23, 2004
King & Spalding, LLP
191 Peachtree Street, N.E., Atlanta, GA 30303
Office: (404)572-4600/ Fax: 404-572-5145

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

Sheet	1	of	3

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Attorney Docket Number	08841.105052 (PHA 2030)

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U.S. PATENT DOCUMENTS

Examiner Initials *	Cite No. [1]	U.S. Patent Document Number	Kind Code [2] (if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages/Relevant Figures Appear
	AA	5,047,407		Belleau *et al.*	09-10-1991	
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	AO	BAUER, M., *et al.*, "Iodide catalysis of oxidations with dimethyl sulfoxide: a convenient two-step synthesis of α-diketones from α-methylene ketones," *J. Org. Chem.*, 40(13):1990-1992 (1975).	
	AP	BELLEAU, B., *et al.*, "Design and activity of a novel class of nucleoside analogs effective against HIV-1," *5th Int. Conf. on AIDS*, Montreal, Canada; June 4-9, 1989; Abstr. No. T.C.O.1. and Poster No. 4576.	

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	BA	CHOI, W.-B., *et al.*, "In situ complexation directs the stereochemistry of N-glycosylation in the synthesis of oxathiolanyl and dioxolanyl nucleoside analogues," *J. Am. Chem. Soc.*, 113(24):9377-9379 (1991).	
	BB	CHOU, T.-S., *et al.*, "A cyclization approach toward five-membered heteroaromatic o-quinodimethanes via fused-3-sulfolenes," *J. Chinese Chem. Soc.*, 44:299-307 (1997).	
	BC	CORBET, A.H., *et al.*, "DAPD," *Curr. Opin. Investig. Drugs*, 2(9):348-353 (2001).	
	BD	EVANS, C.A. *et al.*, "Divergent asymmetric syntheses of dioxolane nucleoside analogues," *Tetrahedron: Asymmetry*, 4(11):2319-2322 (1993).	
	BE	GRESE, T. A., *et al.*, "General approach to halogenated tetrahydrofuran natural products from red algae of the genus *Laurencia*. Total synthesis of (±)-kumausallene and (±)-1-*epi*-kumausallene," *J. Org. Chem.*, 58(9):2468-2477 (1993).	
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	BI	HANESSIAN, S., *et al.*, "Oxidation of alcohols with *N*-halosuccinimides -- new and efficient variants," *Synthesis*, 1981:394-396 (May 1981).	
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	BN	KORNBLUM, N., *et al.*, "A new and selective method of oxidation. The conversion of alkyl halides and alkyl tosylates to aldehydes," *J. Am. Chem. Soc.*, 81:4113-4114 (August 5, 1959).	
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	CA	MARSHALL, J. A., *et al.*, "Stereoselective total synthesis of the pseudopterolide kallolide A," *J. Org. Chem.*, 63(17):5962-5970 (1998).	
	CB	MEWSHAW, J.P., *et al.*, "Dioxolane guanosine, the active form of the prodrug diaminopurine dioxolane, is a potent inhibitor of drug-resistant HIV-1 isolates from patients for whom standard nucleoside therapy fails," *J. Acquir. Immune Defic. Syndr.*, 29(1):11-20 (January 1, 2002).	
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	CD	OHLE, H., "Die Benzoylierung des Erythrits and Darstellung von Derivaten des *O*-Benzoyl-glykolaldehyds," *Chem. Ber.*, 74(2):291-294 (1941).	
	CE	SHIAO, M.-J., *et al.*, "A convenient synthesis of protected α-hydroxyacetaldehydes," *Synthetic Commun.*, 18(4):359-366 (1988).	
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